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The following <u>Listing of the Claims</u> will replace all prior versions and all prior listings of the claims in the present application:

Listing of The Claims:

- (Currently Amended) A method for treating cancer, which comprises administering to a
 mammal in need of such treatment an effective amount of DMXAA or a
 pharmaceutically acceptable salt or ester thereof and administering an effective amount
 of at least one of a compound selected from platinum compounds, vinca alkaloids,
 alkylating agents cyclophosphamide, anthracyclines, topoisomerase I inhibitors,
 antimetabolites and topoisomerase II inhibitors.
- (Currently Amended) A method according to claim 1 for treating cancer, which comprises administering to a mammal in need of such treatment an effective amount of DMXAA or a pharmaceutically acceptable salt or ester thereof and administering an effective amount of at least one of a compound selected from platinum compounds, vinca alkaloids, alkylating agents, anthracyclines, topoisomerase I inhibitors, antimetabolites and topoisomerase II inhibitors, wherein the DMXAA or pharmaceutically acceptable salt or ester thereof and the at least one of a compound selected from platinum compounds, vinca alkaloids, alkylating agents, anthracyclines, topoisomerase I inhibitors, antimetabolites and topoisomerase II inhibitors are administered in a potentiating ratio.
- 3. (Currently Amended) A method according to claim 1 wherein the DMXAA or pharmaceutically acceptable salt or ester thereof and the at least one of a compound selected from platinum compounds, vinca alkaloids, alkylating agents cyclophosphamide, anthracyclines, topoisomerase I inhibitors, antimetabolites and topoisomerase II inhibitors are administered concomitantly.
- 4. (Currently Amended) A method according to claim 1 for treating cancer, which comprises administering to a mammal in need of such treatment an effective amount of DMXAA or a pharmaceutically acceptable salt or ester thereof and administering an effective amount of at least one of a compound selected from platinum compounds, vinca alkaloids, alkylating agents, anthracyclines, topoisomerase I inhibitors, antimetabolites

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and topoisomerase II inhibitors, wherein the DMXAA or pharmaceutically acceptable salt or ester thereof and the at least one of a compound selected from platinum compounds, vinca alkaloids, alkylating agents, anthracyclines, topoisomerase I inhibitors, antimetabolites and topoisomerase II inhibitors are administered sequentially.

- 5. (Currently Amended) A method according to claim 1 for treating cancer, which comprises administering to a mammal in need of such treatment an effective amount of DMXAA or a pharmaceutically acceptable salt or ester thereof and administering an effective amount of at least one of a compound selected from platinum compounds, vinca alkaloids, alkylating agents, anthracyclines, topoisomerase I inhibitors, antimetabolites and topoisomerase II inhibitors, wherein the compound selected from platinum compounds, vinca alkaloids, alkylating agents, anthracyclines, topoisomerase I inhibitors, antimetabolites and topoisomerase II inhibitors is a compound selected from carboplatin, gemcitabine, cisplatin, 5-fluorouracil, cyclophosphamide, etoposide, vincristine, doxorubicin and irinotecan.
- 6. (Original) A method according to claim 5 wherein the compound selected from platinum compounds, vinca alkaloids, alkylating agents, anthracyclines, topoisomerase I inhibitors, antimetabolites and topoisomerase II inhibitors is a compound selected from carboplatin, gemcitabine, cisplatin, 5-fluorouracil, cyclophosphamide, etoposide, vincristine and doxorubicin.
- 7. (Original) A composition comprising a combination of DMXAA or a pharmaceutically acceptable salt or ester thereof and at least one of a compound selected from platinum compounds, vinca alkaloids, alkylating agents, anthracyclines, topoisomerase I inhibitors, antimetabolites and topoisomerase II inhibitors.
- 8. (Original) A composition according to claim 7 wherein the DMXAA or a pharmaceutically acceptable salt or ester thereof and the at least one of a compound selected from platinum compounds, vinca alkaloids, alkylating agents, anthracyclines, topoisomerase I inhibitors, antimetabolites and topoisomerase II inhibitors are present in a potentiating ratio.

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9. (Original) A composition according to claim 7 or 8 wherein the compound selected from platinum compounds, vinca alkaloids, alkylating agents, anthracyclines, topoisomerase I inhibitors, antimetabolites and topoisomerase II inhibitors is a compound selected from carboplatin, gemcitabine, cisplatin, 5-fluorouracil, cyclophosphamide, etoposide, vincristine, doxorubicin and irinotecan.

- 10. (Original) A composition according to claim 9 wherein the compound selected from platinum compounds, vinca alkaloids, alkylating agents, anthracyclines, topoisomerase I inhibitors, antimetabolites and topoisomerase II inhibitors is a compound selected from carboplatin, gemcitabine, cisplatin, 5-fluorouracil, cyclophosphamide, etoposide, vincristine and doxorubicin.
- 11. (Original) A pharmaceutical formulation comprising a combination of DMXAA or a pharmaceutically acceptable salt or ester thereof and at least one of a compound selected from platinum compounds, vinca alkaloids, alkylating agents, anthracyclines, topoisomerase I inhibitors, antimetabolites and topoisomerase II inhibitors in association with one or more pharmaceutically acceptable carriers therefor.
- 12. (Original) A pharmaceutical formulation according to claim 11 wherein the formulation is adapted for intravenous administration.
- 13. (Original) A pharmaceutical formulation according to claim 11 or 12 wherein the DMXAA or pharmaceutically acceptable salt or ester thereof and the at least one of a compound selected from platinum compounds, vinca alkaloids, alkylating agents, anthracyclines, topoisomerase I inhibitors, antimetabolites and topoisomerase II inhibitors are present in a potentiating ratio.
- 14. (Original) A pharmaceutical formulation according to claim 13 wherein the compound selected from platinum compounds, vinca alkaloids, alkylating agents, anthracyclines, topoisomerase I inhibitors, antimetabolites and topoisomerase II inhibitors is a compound selected from carboplatin, gemcitabine, cisplatin, 5-fluorouracil, cyclophosphamide, etoposide, vincristine, doxorubicin and irinotecan.

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15. (Original) A pharmaceutical formulation according to claim 14 wherein the compound selected from platinum compounds, vinca alkaloids, alkylating agents, anthracyclines, topoisomerase I inhibitors, antimetabolites and topoisomerase II inhibitors is a compound selected from carboplatin, gemcitabine, cisplatin, 5-fluorouracil, cyclophosphamide, etoposide, vincristine and doxorubicin.

- 16. (Original) A process for the preparation of a pharmaceutical formulation which process comprises bringing into association a combination of DMXAA or a pharmaceutically acceptable salt or ester thereof and at least one of a compound selected from platinum compounds, vinca alkaloids, alkylating agents, anthracyclines, topoisomerase I inhibitors, antimetabolites and topoisomerase II inhibitors with one or more pharmaceutically acceptable carriers therefor.
- 17. (Original) A process according to claim 16 wherein the DMXAA or pharmaceutically acceptable salt or ester thereof and the at least one of a compound selected from platinum compounds, vinca alkaloids, alkylating agents, anthracyclines, topoisomerase I inhibitors, antimetabolites and topoisomerase II inhibitors are present in a potentiating ratio.
- 18. (Original) A process according to claim 16 or 17 wherein the compound selected from platinum compounds, vinca alkaloids, alkylating agents, anthracyclines, topoisomerase I inhibitors, antimetabolites and topoisomerase II inhibitors is a compound selected from carboplatin, gemcitabine, cisplatin, 5-fluorouracil, cyclophosphamide, etoposide, vincristine, doxorubicin and irinotecan.
- 19. (Original) A process according to claim 18 wherein the compound selected from platinum compounds, vinca alkaloids, alkylating agents, anthracyclines, topoisomerase I inhibitors, antimetabolites and topoisomerase II inhibitors is a compound selected from carboplatin, gemcitabine, cisplatin, 5-fluorouracil, cyclophosphamide, etoposide, vincristine and doxorubicin.
- 20. (Original) A kit comprising in association for separate administration DMXAA or a pharmaceutically acceptable salt or ester thereof and at least one of a compound selected from platinum compounds, vinca alkaloids, alkylating agents, anthracyclines, topoisomerase I inhibitors, antimetabolites and topoisomerase II inhibitors.

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21. (Original) A kit according to claim 20 wherein the DMXAA or pharmaceutically acceptable salt or ester thereof and the at least one of a compound selected from platinum compounds, vinca alkaloids, alkylating agents, anthracyclines, topoisomerase I inhibitors, antimetabolites and topoisomerase II inhibitors are present in a potentiating ratio.

- 22. (Original) A kit according to claim 20 or 21 wherein the compound selected from platinum compounds, vinca alkaloids, alkylating agents, anthracyclines, topoisomerase I inhibitors, antimetabolites and topoisomerase II inhibitors is a compound selected from carboplatin, gemcitabine, cisplatin, 5-fluorouracil, cyclophosphamide, etoposide, vincristine, doxorubicin and irinotecan.
- 23. (Original) A kit according to claim 22 wherein the compound selected from platinum compounds, vinca alkaloids, alkylating agents, anthracyclines, topoisomerase I inhibitors, antimetabolites and topoisomerase II inhibitors is a compound selected from carboplatin, gemcitabine, cisplatin, 5-fluorouracil, cyclophosphamide, etoposide, vincristine and doxorubicin.